## **AMENDMENTS TO THE CLAIMS**

- (Currently amended) Oral A chewing gum pharmaceutical composition comprising containing at least two separate formulations:
  - <u>a core comprising</u> a first formulation, which contains an active compound selected from 2-[4-(diphenylmethyl)-1-piperazinyl]-acetic acids and their amides having the general formula I

#### Formula I

wherein

R<sub>1</sub> is a -COOH group or a -CONH<sub>2</sub> group, and

X<sub>1</sub> and X<sub>2</sub>, taken separately, each represent a hydrogen atom, a halogen atom, a straight-chain or branched C<sub>1</sub>-C<sub>4</sub> alkoxy group or a trifluoromethyl group as well as their pharmaceutically acceptable salts, geometrical isomers, enantiomers, diastereomers and mixtures thereof, and which first formulation does not contain polyols having a molecular weight of less than 300 in a molar ratio between the polyol and active compound of formula I above 10, and

### additionally containing a gum base; and

- <u>a coating comprising</u> a second formulation, which contains one or more solid polyol(s) with a molecular weight of less than 3000 and is free of a compound of formula I, wherein a polyol in the second formulation is a polysaccharide.
- (Original) A composition according to claim 1 wherein the first formulation does
  not contain polyols having a molecular weight of less than 950 in a molar ratio
  between polyol and active compound of formula I above 10, with the exception of
  lactose.
- 3. (Original) A composition according to claim 1 wherein the first formulation does not contain polyols having a molecular weight of less than 300 in a molar ratio between polyol and active compound of formula I above 5.

# 4. (Cancelled)

- 5. (Previously presented) A composition according to claim 1 wherein the first formulation does not contain polyols having a molecular weight of less than 950 in a molar ratio between polyol and active compound of formula I above 5, with the exception of lactose.
- 6. (Previously presented) A composition according to claim 1 wherein the first formulation does not contain polyols having a molecular weight of less than 950, with the exception of lactose.
- 7. **(Previously presented)** A composition according to claim 1 wherein the formulations are prepared in the form of powders, granules, solutions or suspensions.

8.	(Currently amended) A composition according to claim 1 wherein the <u>a</u> polyol in the second formulation is mannitol.
9.	(Cancelled)
10.	(Previously presented) A composition according to claim 1 wherein at least one of the formulations further contains an alkalinizing agent.
11.	(Previously presented) A composition according to claim 10 wherein the alkalinizing agent is sodium citrate.
12.	(Previously presented) A composition according to claim 1 wherein the first formulation further contains one or more excipients selected from cyclodextrins, colloidal anhydrous silica, microcristalline cellulose, magnesium stearate, flavors or colorants.
13.	(Previously presented) A composition according to claim 1 wherein the first formulation further contains non-polyol sweetening agents such as accesulfame K, aspartame, saccharine, saccharine sodium or cyclamate.
14.	(Previously presented) A composition according to claim 1 wherein the active compound in the first formulation is cetirizine dihydrochloride, levocetirizine dihydrochloride or efletirizine dihydrochloride.
15.	(Cancelled)
16.	(Cancelled)
17.	(Cancelled)
18.	(Cancelled)

- 19. (Cancelled)
- 20. (Cancelled)
- 21. (Cancelled)
- 22. (Cancelled)
- 23. (New) A chewing gum pharmaceutical composition comprising:
  - a coating comprising a first formulation, which contains an active compound selected from 2-[4-(diphenylmethyl)-1-piperazinyl]-acetic acids and their amides having the general formula I

#### Formula I

wherein

R<sub>1</sub> is a -COOH group or a -CONH<sub>2</sub> group, and

 $X_1$  and  $X_2$ , taken separately, each represent a hydrogen atom, a halogen atom, a straight-chain or branched  $C_1$ - $C_4$  alkoxy group or a trifluoromethyl group as well as

their pharmaceutically acceptable salts, geometrical isomers, enantiomers, diastereomers and mixtures thereof, and which first formulation does not contain polyols having a molecular weight of less than 300 in a molar ratio between the polyol and active compound of formula I above 10; and

- a core comprising a second formulation, which contains one or more solid polyol(s) with a molecular weight of less than 3000 and is free of a compound of formula I, wherein a polyol in the second formulation is a polysaccharide, and additionally containing a gum base.
- 24. (New) A composition according to claim 23 wherein the first formulation does not contain polyols having a molecular weight of less than 950 in a molar ratio between polyol and active compound of formula I above 10, with the exception of lactose.
- 25. (New) A composition according to claim 23 wherein the first formulation does not contain polyols having a molecular weight of less than 300 in a molar ratio between polyol and active compound of formula I above 5.
- 26. (New) A composition according to claim 23 wherein the first formulation does not contain polyols having a molecular weight of less than 950 in a molar ratio between polyol and active compound of formula I above 5, with the exception of lactose.
- 27. (New) A composition according to claim 23 wherein the first formulation does not contain polyols having a molecular weight of less than 950, with the exception of lactose.
- 28. (New) A composition according to claim 23 wherein the formulations are prepared in the form of powders, granules, solutions or suspensions.

- 29. (New) A composition according to claim 23 wherein a polyol in the second formulation is mannitol.
- 30. (New) A composition according to claim 23 wherein at least one of the formulations further contains an alkalinizing agent.
- 31. (New) A composition according to claim 30 wherein the alkalinizing agent is sodium citrate.
- 32. **(New)** A composition according to claim 23 wherein the first formulation further contains one or more excipients selected from cyclodextrins, colloidal anhydrous silica, microcristalline cellulose, magnesium stearate, flavors or colorants.
- 33. (New) A composition according to claim 23 wherein the first formulation further contains non-polyol sweetening agents such as accesulfame K, aspartame, saccharine, saccharine sodium or cyclamate.
- 34. **(New)** A composition according to claim 23 wherein the active compound in the first formulation is cetirizine dihydrochloride, levocetirizine dihydrochloride or efletirizine dihydrochloride.
- 35. **(New)** A composition according to claim 12 wherein the first formulation further contains a cyclodextrin.
- 36. (New) A composition according to claim 35 wherein the cyclodextrin is beta cyclodextrin.
- 37. (New) A composition according to claim 32 wherein the first formulation further contains a cyclodextrin.

38. (New) A composition according to claim 37 wherein the cyclodextrin is beta cyclodextrin.